

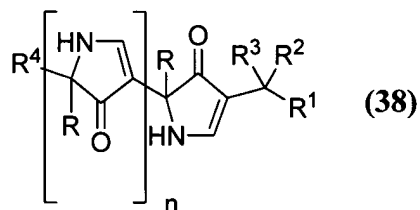
Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

We claim:

1. (Currently amended): A process for preparing a polypyrrolinone having the formula (38):



wherein:

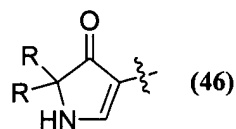
R is independently selected from a group consisting of a straight C₁-C₆ alkyl, a branched C₃-C₇ alkyl, C₃-C₇ cycloalkyl, a straight C₁-C₆ alkenyl, a branched C₃-C₇ alkenyl, C₁-C₄ hydroxyalkyl, C₁-C₄ thioalkyl, C₁-C₄ methylthioalkyl, -(CH₂)_oN(R⁵)₂, -(CH₂)_oCO₂H, -(CH₂)_oCON(R⁵)₂, heteroaryl, phenyl optionally substituted with one to three hydroxyl, ~~lower~~ C₁-C₈ alkoxy, halo, nitro, or cyano groups, and C₇-C₁₂ benzyl optionally substituted with ~~the same groups as above or heteroaryl~~ one to three hydroxyl, C₁-C₈ alkoxy, halo, nitro or cyano groups;

R¹ is hydrogen, hydroxyl, ~~lower~~ C₁-C₈ alkoxy, amino or alkoxycarbonyl-protected amino;

R² is R, carboxyl, a carbonyl linked to a solid support or alkoxycarbonyl;

R³ is R or hydrogen;

R⁴ is R or (46);



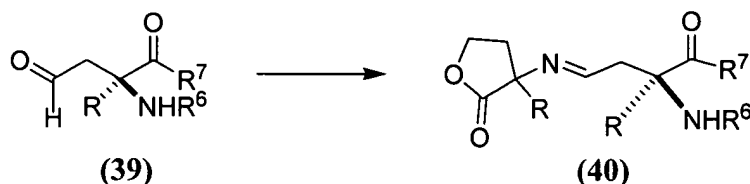
R^5 is hydrogen or ~~lower~~ C_1 - C_8 alkyl;

n is 0 to 3;

o is 1 to 4;

comprising the steps:

- (a) ~~exposing~~ reacting an α -amino- α -substituted-1,4-dioxo compound (39), optionally with an alkoxycarbonyl protecting group, ~~to a plurality of treatments~~ with a 2-substituted-2-aminovalerolactone, trimethylorthoformate, optionally in the presence of a solvent, to produce imine (40)



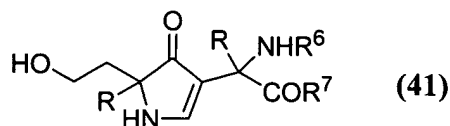
wherein:

R^6 is an amino protecting group,

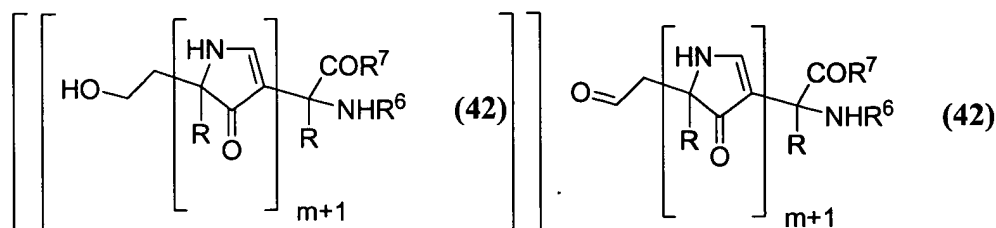
R^7 is a C_1 - C_4 alkoxy or a carboxyl or carbamido linked to a solid support, or

R^6 and R^7 together form a pyrrolinone ring;

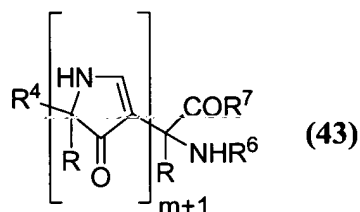
- (b) cyclizing (40) by forming metalloimine carbanion with base optionally in the presence of a crown ether to form a pyrrolinone (41);



- (c) oxidizing the primary alcohol to the corresponding aldehyde;
- (d) repeating steps (a)-(c) m times to produce polypyrrolinone (42);

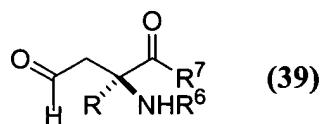


(e) terminating the synthesis by repeating steps (a) through ~~(e)~~(b) using α -substituted amino acid ester in



(f) place of the valerolactone in step ~~(b)~~(a) to yield (43).

- (Original): A process according to claim 1 wherein said polypyrrolinones are substantially diastereomerically pure.
- (Currently amended): A process according to Claim 1 wherein the initial α -amino- α -substituted-1,4-dioxo compound is a compound (39) and R^6 is an alkoxy carbonyl

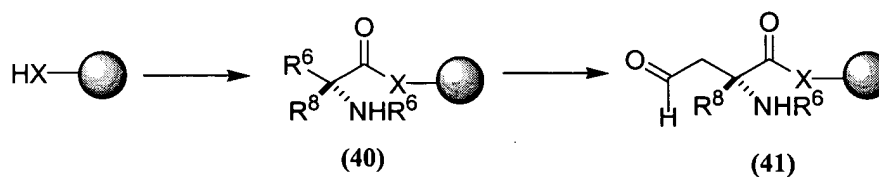


protecting group, R is as defined above and R^7 is a lower C₁-C₈ alkoxy group,

- (Original): A process according to claim 1 wherein the oxidant in step (c) is oxalyl chloride, a tertiary amine and DMSO.
- (Original): A process according to Claim 4 wherein the tertiary amine is DBU or di-*iso*-propylethyl amine.

6. (Original): A process according to Claim 1 wherein the crown ether in step (b) is 18-crown-6.
7. (Original): A process according to Claim 1 wherein the base in step (b) is potassium hexamethyldisilazane.
8. (Withdrawn): A solid-phase process according to claim 1 wherein R^7 is a carboxyl or carbamido linked to a solid support further comprising the steps of:

(f) attaching a latent aldehyde (40) to a solid support wherein and converting the latent aldehyde to an aldehyde (41);

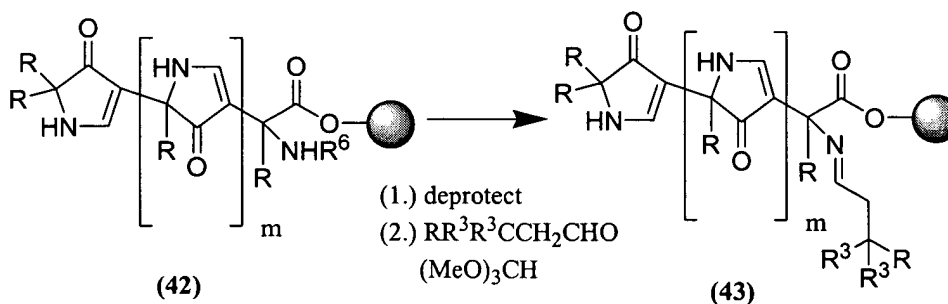


wherein:

R^8 is 3-methyl-1-but-2-enyl, 2,2-dimethoxyethyl, 2-hydroxyethyl, and

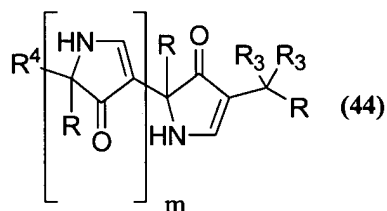
X is nitrogen or oxygen;

(g) repeating steps (a)-(c) m times and terminating the synthesis as in step (e) to produce polypyrrolinone (42);

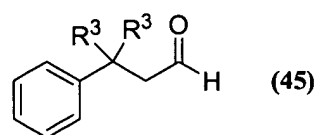


(h) cleaving the polypyrrolinone from the resin by deprotecting the α -amino group, and exposing the α -amino acid to a plurality of treatments with an aldehyde, trimethylorthoformate, optionally in the presence of a solvent, to produce the corresponding imine (43); and,

(i) cyclizing (43) by forming the metalloimine carbanion with base, optionally in the presence of a crown ether, to produce a pyrrolinone (44).



9. (Withdrawn): A process according to claim 7 wherein the oxidant in step (c) is oxalyl chloride, a tertiary amine and DMSO.
10. (Withdrawn): A process according to Claim 7 wherein the tertiary amine is DBU or di-*iso*-propylethyl amine.
11. (Withdrawn): A process according to Claim 7 wherein the crown ether in step (b) is 18-crown-6.
12. (Withdrawn): A process according to Claim 7 wherein the base in step (b) is potassium hexamethyldisilazane.
13. (Withdrawn): A process according to Claim 7 wherein R⁶ is a trialkylsilylethoxycarbonyl group.
14. (Withdrawn): A process according to Claim 7 wherein the aldehyde in step (h) is a 3-phenylpropionaldehyde (45) derivative optionally substituted at the 3-position with one or two R³ substituents.



15. (Withdrawn): A process according to Claim 7 wherein the aldehyde in step (h) is 3-phenylpropionaldehyde.